DEFENSE

Cooperative Research Projects

Agreement Between the UNITED STATES OF AMERICA and SINGAPORE

Signed at Singapore and Washington July 20 and August 14, 2009



NOTE BY THE DEPARTMENT OF STATE

Pursuant to Public Law 89—497, approved July 8, 1966 (80 Stat. 271; 1 U.S.C. 113)—

"...the Treaties and Other International Acts Series issued under the authority of the Secretary of State shall be competent evidence... of the treaties, international agreements other than treaties, and proclamations by the President of such treaties and international agreements other than treaties, as the case may be, therein contained, in all the courts of law and equity and of maritime jurisdiction, and in all the tribunals and public offices of the United States, and of the several States, without any further proof or authentication thereof."

SINGAPORE

Defense: Cooperative Research Projects

Agreement signed at Singapore and Washington July 20 and August 14, 2009; Entered into force August 14, 2009.

AGREEMENT BETWEEN THE

DEPARTMENT OF DEFENSE OF THE UNITED STATES OF AMERICA

AND THE

MINISTRY OF DEFENSE OF THE REPUBLIC OF SINGAPORE

RELATING TO COOPERATIVE RESEARCH PROJECTS FOR SURVEILLANCE OF

MALARIA DRUG RESISTANCE

ARTICLE I INTRODUCTION

The Department of Defense of the United States of America (U.S. DoD) and the Ministry of Defense of the Republic of Singapore (MINDEF), pursuant to the Strategic Framework Agreement Between the United States of America and the Republic of Singapore for a Closer Cooperation Partnership in Defense and Security, signed in Washington DC, on 12 July 2005, agree to establish Surveillance for Malaria Drug Resistance as a cooperative medical research project. The terms of the Strategic Framework Agreement between the United States of America and the Republic of Singapore for a Closer Cooperation Partnership in Defense and Security Agreement shall govern the execution of this Agreement.

ARTICLE II DEFINITION OF ABBREVIATIONS

DMERI Defence Medical and Environmental Research Institute, a research entity

of DSO

DSO DSO National Laboratories, Singapore

DSTA Defence Science and Technology Agency, a statutory board of

MINDEF, Singapore

MINDEF Ministry of Defence, Singapore

HQMC Headquarters Medical Corps, Singapore Armed Forces

SAF Singapore Armed Forces
DoD Department of Defense, USA

NAMRU-2 Naval Medical Research Unit Two, a research entity of DoD NMRC Navy Medical Research Center, a research entity of DoD

GEIS Global Emerging Infectious Disease Surveillance and Responses, DoD

USN United States Navy

RDT&E Research, Development, Testing, and Evaluation Project

ARTICLE III OBJECTIVES

The objective of this Agreement is to facilitate cooperation between the U. S. DoD and the MINDEF of the Republic of Singapore to conduct collaborative biomedical RDT&E projects on malaria drug resistance. The specific objectives of this Agreement are:

- 1. To establish a joint surveillance network to determine the species of malaria parasite and the drug resistance profile of imported infections in Singapore.
- 2. To conduct molecular marker testing on malaria parasites imported from different geographical regions to monitor the polymorphisms associated with antimalarial drug resistance.
- 3. To conduct in vitro drug susceptility testing on malaria parasites imported from different geographical regions to monitor intrinsic parasite drug sensitivity (ED50).

ARTICLE IV SCOPE OF WORK

The following work shall be performed under this Agreement:

- 1. Phase 1 will be to identify and establish collaboration with hospitals and physicians in Singapore to participate in the malaria surveillance project and to establish laboratory capability, standard operating procedures (SOPs) and protocols, and monthly reporting formats.
- 2. Phase 2 will be to conduct training of laboratory personnel to perform laboratory assays according to laboratory SOP and to establish a laboratory proficiency programme with participating US DoD laboratories.
- 3. Phase 3 will be daily operations of the joint surveillance network and providing annual reports on the activities, performance and achievements of the joint surveillance network. The collection of specimens for this project will be governed by Institutional Review Board (IRB) protocol approved by Singapore and reviewed by the U.S.

ARTICLE V. SHARING OF TASKS

The sharing of tasks shall be as follows:

The MINDEF agrees to provide the following:

- 1. Review, submit and obtain IRB approval. Provide approval documentation to the U.S. for review.
- 2. Collect clinical and epidemiological and specimen from patients with confirmed malaria diagnosis.
- 3. Provide laboratory personnel for training and performance of laboratory assays and procedures.
- 4. Participate in the laboratory proficiency program with participating US DoD laboratories.

The U.S. DoD agrees to provide the following:

- 1. Review the IRB protocol in accordance with SECNAVINST 3900.39D.
- 2. Provide standardized laboratory assays and protocols used by DoD laboratories for identification, molecular genotyping and in vitro drug susceptility testing for characterization of malaria parasites.
- 3. Provide laboratory personnel with hands-on training on the conduct and performance of designated laboratory assays and protocols.
- 4. Provide standardized antimalarial drugs and detection capability for performance of in vitro drug resistance testing using established fluorescent detection methods.

The MINDEF and the DoD agree to jointly provide the following:

- 1. Identity laboratory assays and protocols to support implementation of the objectives of this project and identify job functions for DMERI, DSTA, and NAMRU-2 personnel.
- 2. Assist in establishing laboratory capability at DMERI.
- 3. Preparation of reporting procedures and generation of annual reports on the activities, performance and achievements of the joint surveillance network.

ARTICLE VI BREAKDOWN AND SCHEDULE OF TASKS

1.	The project s	shall proce	ed according to t	he fol	lowing	concurrent	phases a	nd sc	hed	ule
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2. Phase 1

Duration

a. Establish project management responsibilities and timelines

3 months

- (1) Complete study proposal and establish collaboration with hospitals and clinics.
- (2) Update and verify job functions for DMERI, NAMRU-2, and hospital investigators and staff.
- (3) Develop IRB Protocol.
- 3. Phase II.
 - a. Establish laboratory support capabilities

3 months

- (1) Obtain IRB Approval and recruit subjects.
- (2) Procure reagents and equipment.
- (3) Establish laboratory protocols for blood collection, shipping and delivery to the laboratory, microscopy, molecular marker testing, and in vitro drug sensitivity tests.
- (4) Conduct laboratory hands-on training of laboratory assays and analysis.
- (5) Establish reporting procedures and laboratory proficiency programme.
- 3. Phase III

30 months

- a. Initiate the study in accordance with the approved IRB protocol and maintain laboratory operations and reporting procedures.
 - (1) Coordinate and execute collection of samples, laboratory activities and reporting, and participation in proficiency programme.
 - (2) Provide annual review reports.
- 4. The project is expected to be completed 31 months from the date of commencement 1.

ARTICLE VII MANAGEMENT

1. This Agreement shall be directed and administered on behalf of the Parties by one Project Officer (PO) from each Party. The POs are:

US PO:

Dr. Gary T. Brice, LCDR, MSC, USN

Title/Position:

Head, Department of Immunology Naval Medical Research Unit No. 2

Organization:

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ARTICLE VIII FINANCIAL PROVISIONS

Each Party will pay for its own expenses resulting from efforts undertaken pursuant to this Agreement.

ARTICLE IX CLASSIFICATION

The existence of this Agreement and its contents is UNCLASSIFIED. No Classified Information shall be exchanged under this Agreement

ARTICLE X PRINCIPAL ORGANIZATIONS INVOLVED

- 1. The United States of America: Naval Medical Research Unit No. 2, Singapore Detachment United States Navy
- 2. Republic of Singapore:
 - a. Defence Medical and Environmental Research Institute, DSO National Laboratories

ARTICLE XI

ENTRY INTO FORCE, DURATION AND TERMINATION

1.	This Agreement shall enter into force upon signature by the Agreement Managemen	t
Agents	, and shall remain in force for three years unless terminated by either Party. It may b	e
extende	ed by written agreement of the Parties.	

2. IN WITNESS WHEREOF the undersigned, duly authorized by their respective Governments, have signed this Agreement.										
Done in duplicate, this 14th day of August 200	9									
For the U.S. MA	For Singapore MA									
1. M. Rommon, Sp. Signature	Signature									
A. M. ROBINSON, JR, VADM, MC, USN Name	TAN PENG YAM Name									
Surgeon General of the Navy Title	Deputy Chief Executive (Operations) Defence Science & Technology Agency Title									
14 August 2009 Date	20 July 2009 Date									
United States of America Location	Republic of Singapore Location									